Agency for Toxic Substances and Disease Registry [60Day-21-0048; Docket No. ATSDR-2021-0007]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Agency for Toxic Substances and Disease Registry (ATSDR), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Agency for Toxic Substances and Disease Registry (ATSDR), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled ATSDR Exposure

Investigations (EIs). The information collection is designed to evaluate public health issues at a site resulting from environmental exposure. ATSDR EIs fill data gaps by conducting environmental and biological sampling.

DATES: ATSDR must receive written comments on or before [INSERT DATE 60 DAYS AFTER PUBLICATION DATE IN THE FEDERAL REGISTER].

ADDRESSES: You may submit comments, identified by Docket No.

ATSDR-2021-0007 by any of the following methods:

- Federal eRulemaking Portal: Regulations.gov. Follow the instructions for submitting comments.
- Mail: Jeffrey M. Zirger, Information Collection Review
  Office, Centers for Disease Control and Prevention, 1600
  Clifton Road, N.E., MS-D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency
name and Docket Number. ATSDR will post, without change, all
relevant comments to Regulations.gov.

Please note: Submit all comments through the Federal eRulemaking portal (regulations.gov) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road, N.E., MS-D74, Atlanta, Georgia 30329; phone: 404-639-7118; E-mail: omb@cdc.gov.

## SUPPLEMENTARY INFORMATION:

Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501-3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the Federal Register concerning each proposed collection of information, including each new proposed collection, each

proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is

necessary for the proper performance of the functions of the

agency, including whether the information will have practical

utility;

- 2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- 3. Enhance the quality, utility, and clarity of the information to be collected;
- 4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses; and
- 5. Assess information collection costs.

Proposed Project

ATSDR Exposure Investigations (EIs) (OMB Control No. 0923-0048, Exp. 04/30/2022) - Extension - Agency for Toxic Substances and Disease Registry (ATSDR).

## Background and Brief Description

The Agency for Toxic Substances and Disease Registry

(ATSDR) is requesting a three-year extension of this generic clearance to allow the agency to conduct exposure investigations (EIs), through methods developed by ATSDR. After a chemical release or suspected release into the environment, EIs are usually requested by officials of a state health agency, county health department, the Environmental Protection Agency (EPA), the general public, and/or ATSDR staff. EI results are used by public health professionals, environmental risk managers, and other decision makers to determine if current conditions warrant intervention strategies to minimize or eliminate human exposure.

All of ATSDR's targeted biological assessments (e.g., urine, blood) and some of the environmental investigations (e.g., air, water, soil, dust, or food sampling) involve participants to determine whether they are or have been exposed to unusual levels of pollutants at specific locations (e.g., where people live, spend leisure time, or anywhere they might come into contact with contaminants under investigation).

Questionnaires, appropriate to the specific contaminant, are generally needed in about half of the EIs (at most, approximately 12 per year) to assist in interpreting the biological or environmental sampling results. ATSDR collects contact information (e.g., name, address, phone number) to provide the participant with their individual results. ATSDR also collects information on other possible confounding sources of chemical(s) exposure such as medicines taken, foods eaten, hobbies, jobs, etc. In addition, ATSDR asks questions on recreational or occupational activities that could increase a participant's exposure potential. The information collected represents an individual's exposure history.

The number of questions can vary depending on the number of chemicals being investigated, the route of exposure (e.g., breathing, eating, touching), and the number of other sources of the chemical(s) (e.g., products used, jobs). We use approximately 12-20 questions about the pertinent environmental exposures per investigation. A question bank is available for health assessors to use as a basis of questions to be asked during the EI, but EI-specific questions may be included as appropriate.

Typically, the number of participants in an individual EI ranges from 10 to 100. Participation is

completely voluntary, and there are no costs to participants other than their time. Based on a maximum of 12 EIs per year and 100 participants each, the estimated annualized burden hours are 600.

Estimated Annualized Burden Hours

| Type of       | Form Name | Number   | Number   | Average  | Total    |
|---------------|-----------|----------|----------|----------|----------|
| Respondent    |           | of       | of       | Burden   | Burden   |
|               |           | Responde | Response | per      | (in hr.) |
|               |           | nts      | s per    | Response |          |
|               |           |          | Responde | (in hr.) |          |
|               |           |          | nt       |          |          |
| Exposure      | Chemical  |          |          |          |          |
| Investigation | Exposure  | 1,200    | 1        | 30/60    | 600      |
| Participants  | Questions |          |          |          |          |
| Total         |           |          |          |          | 600      |

## Jeffrey M. Zirger,

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Centers for Disease Control and Prevention.

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